

Amendments to the Claims:

Please cancel claims 1-48, 50-55, 64-65 and 73-74 without prejudice.

Please add new claims 75-86.

Please amend claims 49, 56-62 and 68-72 as follows:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Claims 1-48 (Cancelled)

49. (Currently amended) A method for diagnosing colon, stomach or prostate cancer comprising: a) determining the expression of one or more genes a gene comprising or encoding a nucleic acid sequence selected from the group consisting of ~~SEQ ID NO:606, SEQ ID NO:4,~~ ~~SEQ ID NO:10, SEQ ID NO:26, SEQ ID NO:32, SEQ ID NO:40, SEQ ID NO:46, SEQ ID NO:54, SEQ ID NO:60, SEQ ID NO:68, SEQ ID NO:76, SEQ ID NO:86, SEQ ID NO:96, SEQ ID NO:102, SEQ ID NO:114, SEQ ID NO:122, SEQ ID NO:128, SEQ ID NO:140, SEQ ID NO:143, SEQ ID NO:149, SEQ ID NO:159, SEQ ID NO:165, SEQ ID NO:183, SEQ ID NO:191, SEQ ID NO:199, SEQ ID NO:205, SEQ ID NO:211, SEQ ID NO:219, SEQ ID NO:225, SEQ ID NO:231, SEQ ID NO:242, SEQ ID NO:248, SEQ ID NO:254, SEQ ID NO:266, SEQ ID NO:272, SEQ ID NO:278, SEQ ID NO:284, SEQ ID NO:290, SEQ ID NO:296, SEQ ID NO:308, SEQ ID NO:314, SEQ ID NO:320, SEQ ID NO:328, SEQ ID NO:334, SEQ ID NO:342, SEQ ID NO:350, SEQ ID NO:356, SEQ ID NO:362, SEQ ID NO:368, SEQ ID NO:374, SEQ ID NO:380, SEQ ID NO:390, SEQ ID NO:398, SEQ ID NO:410, SEQ ID NO:416, SEQ ID NO:422, SEQ ID NO:444, SEQ ID NO:450, SEQ ID NO:458, SEQ ID NO:466, SEQ ID NO:474, SEQ ID NO:486, SEQ ID NO:492, SEQ ID NO:500, SEQ ID NO:508, SEQ ID NO:514, SEQ ID NO:520, SEQ ID NO:526, SEQ ID NO:534, SEQ ID NO:540, SEQ ID NO:554, SEQ ID NO:564, SEQ ID NO:570, SEQ ID NO:576, SEQ ID NO:582, SEQ ID NO:588, SEQ ID NO:594, SEQ ID NO:600, SEQ ID NO:612, SEQ ID NO:620, SEQ ID NO:628, SEQ ID NO:634, SEQ ID NO:640, SEQ ID~~

NO:648, SEQ ID NO:654, SEQ ID NO:664, SEQ ID NO:672, SEQ ID NO:680, SEQ ID NO:692, SEQ ID NO:698, SEQ ID NO:704, SEQ ID NO:710, SEQ ID NO:716, SEQ ID NO:726, SEQ ID NO:732, SEQ ID NO:735, SEQ ID NO:741, SEQ ID NO:747, SEQ ID NO:753, SEQ ID NO:759, SEQ ID NO:775, SEQ ID NO:781, SEQ ID NO:787, SEQ ID NO:795, SEQ ID NO:801, SEQ ID NO:811, SEQ ID NO:817, SEQ ID NO:823, SEQ ID NO:829, SEQ ID NO:835, SEQ ID NO:841, SEQ ID NO:847, SEQ ID NO:853, SEQ ID NO:859, SEQ ID NO:865, SEQ ID NO:871, SEQ ID NO:877, SEQ ID NO:887, SEQ ID NO:899, SEQ ID NO:905, SEQ ID NO:911, SEQ ID NO:919, SEQ ID NO:925, SEQ ID NO:931, SEQ ID NO:937, SEQ ID NO:945, SEQ ID NO:951, SEQ ID NO:957, SEQ ID NO:963, SEQ ID NO:969, SEQ ID NO:975, SEQ ID NO:607, SEQ ID NO:5, SEQ ID NO:11, SEQ ID NO:13, SEQ ID NO:15, SEQ ID NO:17, SEQ ID NO:19, SEQ ID NO:21, SEQ ID NO:27, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:41, SEQ ID NO:47, SEQ ID NO:49, SEQ ID NO:55, SEQ ID NO:61, SEQ ID NO:63, SEQ ID NO:69, SEQ ID NO:71, SEQ ID NO:77, SEQ ID NO:79, SEQ ID NO:81, SEQ ID NO:87, SEQ ID NO:89, SEQ ID NO:91, SEQ ID NO:97, SEQ ID NO:103, SEQ ID NO:105, SEQ ID NO:107, SEQ ID NO:109, SEQ ID NO:115, SEQ ID NO:117, SEQ ID NO:123, SEQ ID NO:129, SEQ ID NO:131, SEQ ID NO:133, SEQ ID NO:135, SEQ ID NO:141, SEQ ID NO:144,

SEQ ID NO:150 , SEQ ID NO:152, and SEQ ID NO:154

, SEQ ID NO:160, SEQ ID NO:166, SEQ ID NO:168, SEQ ID NO:170, SEQ ID NO:172, SEQ ID NO:174, SEQ ID NO:176, SEQ ID NO:184, SEQ ID NO:186, SEQ ID NO:192, SEQ ID NO:194, SEQ ID NO:200, SEQ ID NO:206, SEQ ID NO:212, SEQ ID NO:214, SEQ ID NO:220, SEQ ID NO:226, SEQ ID NO:232, SEQ ID NO:234, SEQ ID NO:243, SEQ ID NO:249, SEQ ID NO:255, SEQ ID NO:257, SEQ ID NO:259, SEQ ID NO:267, SEQ ID NO:273, SEQ ID NO:279, SEQ ID NO:285, SEQ ID NO:291, SEQ ID NO:297, SEQ ID NO:299, SEQ ID NO:301, SEQ ID NO:303, SEQ ID NO:309, SEQ ID NO:315, SEQ ID NO:321, SEQ ID NO:323, SEQ ID NO:329, SEQ ID NO:335, SEQ ID NO:337, SEQ ID NO:343, SEQ ID NO:345, SEQ ID NO:351, SEQ ID NO:357, SEQ ID NO:363, SEQ ID NO:369, SEQ ID NO:375, SEQ ID NO:381, SEQ ID NO:383, SEQ ID NO:385, SEQ ID NO:391, SEQ ID NO:393, SEQ ID NO:399, SEQ ID NO:401, SEQ ID NO:403, SEQ ID NO:405, SEQ ID NO:411, SEQ ID NO:417, SEQ ID NO:423, SEQ ID NO:425, SEQ ID

NO:427, SEQ\_ID NO:429, SEQ\_ID NO:431, SEQ\_ID NO:433, SEQ\_ID NO:435, SEQ\_ID NO:437, SEQ\_ID NO:439, SEQ\_ID NO:445, SEQ\_ID NO:451, SEQ\_ID NO:453, SEQ\_ID NO:459, SEQ\_ID NO:461, SEQ\_ID NO:467, SEQ\_ID NO:469, SEQ\_ID NO:475, SEQ\_ID NO:477, SEQ\_ID NO:479, SEQ\_ID NO:481, SEQ\_ID NO:487, SEQ\_ID NO:493, SEQ\_ID NO:495, SEQ\_ID NO:501, SEQ\_ID NO:509, SEQ\_ID NO:515, SEQ\_ID NO:521, SEQ\_ID NO:527, SEQ\_ID NO:529, SEQ\_ID NO:535, SEQ\_ID NO:541, SEQ\_ID NO:543, SEQ\_ID NO:545, SEQ\_ID NO:547, SEQ\_ID NO:549, SEQ\_ID NO:555, SEQ\_ID NO:557, SEQ\_ID NO:559, SEQ\_ID NO:565, SEQ\_ID NO:571, SEQ\_ID NO:577, SEQ\_ID NO:583, SEQ\_ID NO:589, SEQ\_ID NO:595, SEQ\_ID NO:601, SEQ\_ID NO:613, SEQ\_ID NO:615, SEQ\_ID NO:621, SEQ\_ID NO:623, SEQ\_ID NO:629, SEQ\_ID NO:635, SEQ\_ID NO:641, SEQ\_ID NO:643, SEQ\_ID NO:649, SEQ\_ID NO:655, SEQ\_ID NO:657, SEQ\_ID NO:665, SEQ\_ID NO:667, SEQ\_ID NO:673, SEQ\_ID NO:675, SEQ\_ID NO:681, SEQ\_ID NO:683, SEQ\_ID NO:685, SEQ\_ID NO:687, SEQ\_ID NO:693, SEQ\_ID NO:699, SEQ\_ID NO:705, SEQ\_ID NO:711, SEQ\_ID NO:717, SEQ\_ID NO:719, SEQ\_ID NO:721, SEQ\_ID NO:727, SEQ\_ID NO:733, SEQ\_ID NO:736, SEQ\_ID NO:742, SEQ\_ID NO:748, SEQ\_ID NO:754, SEQ\_ID NO:760, SEQ\_ID NO:762, SEQ\_ID NO:764, SEQ\_ID NO:766, SEQ\_ID NO:768, SEQ\_ID NO:770, SEQ\_ID NO:776, SEQ\_ID NO:782, SEQ\_ID NO:788, SEQ\_ID NO:796, SEQ\_ID NO:802, SEQ\_ID NO:804, SEQ\_ID NO:806, SEQ\_ID NO:812, SEQ\_ID NO:818, SEQ\_ID NO:824, SEQ\_ID NO:830, SEQ\_ID NO:836, SEQ\_ID NO:842, SEQ\_ID NO:848, SEQ\_ID NO:854, SEQ\_ID NO:860, SEQ\_ID NO:866, SEQ\_ID NO:872, SEQ\_ID NO:878, SEQ\_ID NO:880, SEQ\_ID NO:882, SEQ\_ID NO:888, SEQ\_ID NO:890, SEQ\_ID NO:892, SEQ\_ID NO:894, SEQ\_ID NO:900, SEQ\_ID NO:906, SEQ\_ID NO:912, SEQ\_ID NO:914, SEQ\_ID NO:920, SEQ\_ID NO:926, SEQ\_ID NO:932, SEQ\_ID NO:938, SEQ\_ID NO:940, SEQ\_ID NO:946, SEQ\_ID NO:952, SEQ\_ID NO:958, SEQ\_ID NO:964, SEQ\_ID NO:970, SEQ\_ID NO:976, SEQ\_ID NO:978 and SEQ\_ID NO:980 in a first tissue type of a first individual; and b) comparing said expression of said gene(s) gene from a second normal tissue type from said first individual or a second unaffected individual; wherein a difference in said expression indicates that the first individual has colon, stomach or prostate cancer.

56. (Currently amended) A method for diagnosing colon, stomach or prostate cancer comprising comparing a level of proteasome component C7-I mRNA in a patient sample comprising colon, stomach or prostate tissue to the level of the proteasome component C7-I mRNA in a normal control; wherein an increase a difference of at least 50% from between the level in the patient sample relative to the level in the normal control indicates that the patient has or is predisposed to colon, stomach or prostate cancer.

57. (Currently amended) The method of claim 56 wherein the proteasome component C7-I mRNA comprises a nucleotide sequence at least 95% identical to SEQ ID NO:152, said mRNA encoding a polypeptide with threonine endopeptidase activity.

58. (Currently amended) The method of claim 56 wherein the proteasome component C7-I mRNA comprises a nucleotide sequence at least 98% identical to SEQ ID NO:152, said mRNA encoding a polypeptide with threonine endopeptidase activity.

59. (Previously presented) The method of claim 56 wherein the proteasome component C7-I mRNA comprises the nucleotide sequence of SEQ ID NO:152.

60. (Currently amended) The method of claim 56 wherein an increase a difference of at least 100% from between the level of the proteasome component C7-I mRNA in the patient sample relative to the normal control indicates that the patient has or is predisposed to colon, stomach or prostate cancer.

61. (Currently amended) A method for diagnosing colon, stomach or prostate cancer comprising detecting evidence of differential expression of proteasome component C7-I in a patient sample, wherein evidence of differential expression of proteasome component C7-I indicates that the patient has colon, stomach or prostate cancer.

62. **(Currently amended)** The method of claim 61 wherein ~~evidence of differential expression is detected by measuring the level of a proteasome component C7-I expression product.~~

63. **(Previously presented)** The method of claim 62 wherein the expression product is a protein or mRNA.

Claims 64-65 **(Cancelled)**

66. **(Previously presented)** The method of claim 62 wherein the level of a proteasome component C7-I expression product in the patient sample is compared to a control.

67. **(Previously presented)** The method of claim 66 wherein the control comprises normal colon, stomach or prostate tissue.

68. **(Currently amended)** The method of claim 66 wherein the level of the expression product in the patient sample is increased differs by at least 200% relative to the control.

69. **(Currently amended)** The method of claim 61 wherein ~~evidence of differential expression is detected by measuring the level of a proteasome component C7-I expression product~~ said expression product comprising a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of SEQ ID NO:150, SEQ ID NO:152, and SEQ ID NO:154 said expression product encoding a polypeptide with threonine endopeptidase activity.

70. **(Currently amended)** The method of claim 61 wherein ~~evidence of differential expression is detected by measuring the level of a proteasome component C7-I expression product~~ comprising a nucleotide sequence at least 98% identical to SEQ ID NO:152, said expression product encoding a polypeptide with threonine endopeptidase activity.

71. (Currently amended) The method of claim 61 wherein evidence of differential expression is detected by measuring the level of a proteasome component C7-I expression product comprising the nucleotide sequence of SEQ ID NO:152.

72. (Currently amended) A method of diagnosing colon, stomach or prostate cancer in a patient comprising:

(a) contacting a polynucleotide that hybridizes under highly stringent conditions to the complement of a nucleotide sequence selected from the group consisting of comprising SEQ ID NO:150, SEQ ID NO:152 and SEQ ID NO:154 with nucleic acids of a patient colon, stomach or prostate sample under binding conditions suitable to form a duplex, wherein hybridization is performed at 50°C to 60°C in 5 X SSC (0.9 mM saline /0.9 mM sodium citrate); and

(b) comparing the amount of the duplex formed to the amount of duplex formed when the polynucleotide is contacted with nucleic acids of a non-cancerous colon, stomach or prostate control, wherein increased levels of a difference of at least 50% in the amount of duplex formed upon contacting said polynucleotide with said nucleic acids of the patient sample compared to the amount of duplex formed upon contacting said polynucleotide and said with the nucleic acids of the non-cancerous control indicates that the patient has colon, stomach or prostate cancer.

Claims 73-74 (Cancelled)

75. (New) The method of claim 56 wherein the proteasome component C7-I mRNA comprises a nucleotide sequence at least 95% identical to SEQ ID NO:150.

76. (New) The method of claim 56 wherein the proteasome component C7-I mRNA comprises a nucleotide sequence at least 98% identical to SEQ ID NO:150.

77. (New) The method of claim 56 wherein the proteasome component C7-I mRNA comprises SEQ ID NO:150.

78. (New) The method of claim 61 wherein differential expression is detected by measuring the level of a proteasome component C7-I expression product at least 98% identical to SEQ ID NO:150.

79. (New) The method of claim 61 wherein differential expression is detected by measuring the level of a proteasome component C7-I expression product comprising SEQ ID NO:150.

80. (New) The method of claim 56 wherein the proteasome component C7-I mRNA comprises a nucleotide sequence at least 95% identical to SEQ ID NO:154.

81. (New) The method of claim 56 wherein the proteasome component C7-I mRNA comprises a nucleotide sequence at least 98% identical to SEQ ID NO:154.

82. (New) The method of claim 56 wherein the proteasome component C7-I mRNA comprises SEQ ID NO:154.

83. (New) The method of claim 61 wherein differential expression is detected by measuring the level of a proteasome component C7-I expression product at least 98% identical to SEQ ID NO:154.

84. (New) The method of claim 61 wherein differential expression is detected by measuring the level of a proteasome component C7-I expression product comprising SEQ ID NO:154.

85. (New) The method of any one of claims 57 or 69 wherein the expression product encodes a threonine endopeptidase.

86. (New) The method of any one of claims 49, 56, 61 or 72 wherein the cancer is colon cancer.